AMENDMENTS TO THE CLAIMS:

This listing of claims will replace all prior versions, and listings, of claims in the application:

1-167 (cancelled).

168 (previously presented). A method of incorporating a synthetic molecule construct of the structure $F-S_1-S_2-L$ into the lipid bi-layer of a cell or a multi-cellular structure including the step:

of contacting a suspension of the cell or multi-cellular structure with the synthetic molecule construct for a time and at a temperature sufficient to allow incorporation where:

F is a mono-, di-, tri- or oligo- saccharide;

S₁ is 2-aminoethyl, 3-aminopropyl, 4-aminobutyl, or 5-aminopentyl;

 S_2 is $-CO(CH_2)_2CO$ -, $-CO(CH_2)_3CO$ -,

-CO(CH₂)₄CO- or -CO(CH₂)₅CO-; and

L is a diacyl- or dialkyl- glycerophospholipid.

169 (previously presented). The method according to claim 168 where the construct includes the substructure:

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where:

n = 3 to 5and M is H.

170 (previously presented). The method according to claim 168 where the cell or multi-cellular structure is of human or murine origin.

171 (previously presented). The method according to claim 168 where the concentration of the construct in the suspension is in the range 0.1 to 10 mg/mL.

172 (previously presented). The method according to 168 where the suspension of the cell or multi-cellular structure is contacted with the construct at a temperature in the range 2 to 37 °C.

173 (previously presented). The method according to claim 172 where the suspension of the cell or multi-cellular structure is contacted with the construct at a temperature in the range 2 to 25 °C.

174 (previously presented). The method according claim 173 where the suspension of the cell or multi-cellular structure is contacted with the construct at a temperature in the range 2 to 4 °C.

175 (previously presented). The method according to claim 168 where F is selected from the group consisting of GalNAca1-3(Fuca1-2)Galß; Gala1-3Galß; Gala5;

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Galα1-3(Fucα1-2)Galß; NeuAcα2-3Galß; NeuAcα2-6Galß; Fucα1-2Galß; Galß1-4GlcNAcß1-6(Galß1-4GlcNAcß1-3)Galß; Fucα1-2Galß1-4GlcNAcß1-6(Fucα1-2Galß1-4GlcNAcß1-6)Galß1-4GlcNAcß1-3)Galß; Fucα1-2Galß1-4GlcNAcß1-6(NeuAcα2-3Galß1-4GlcNAcß1-3)Galß; NeuAcα2-3Galß1-4GlcNAcß1-6(NeuAcα2-3Galß1-4GlcNAcß1-3)Galß; Galα1-4Galß1-4Glc; GalNAcß1-3Galα1-4Galß1-4Glc; GalNAcß1-3Galα1-4Galß1-4Glc; and GalNAcß1-3GalNAcß1-3Galα1-4Galß1-4Glc.

176 (previously presented). The method according to claim 175 where F is selected from the group consisting of GalNAcα1-3(Fucα1-2)Galß and Galα1-3(Fucα1-2)Galß.

177 (previously presented). The method according to claim 168 where S_1 is 3-aminopropyl.

178 (previously presented). The method according to claim 168 where L is selected from the group consisting of: 1,2-O-dioleoyl-sn-glycero-3-phosphatidylethanolamine (DOPE) and 1,2-O-distearyl-sn-glycero-3-phosphatidylethanolamine (DSPE).

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designated A_{tri}-sp-Ad-DOPE (I).

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designated A_{tri}-sp-Ad-DSPE (III).

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designated B_{tri}-sp-Ad-DOPE (VI).

designated H_{tri}-sp-Ad-DOPE (VII).

designated H_{di}-sp-Ad-DOPE (VIII).

184 (previously presented).

The method according to claim 168 where the

construct is:

designated Galß-sp-Ad-DOPE (IX).

designated Fucα1-2Galβ1-3GlcNAcβ1-3Galβ1-4GlcNAc-sp-Ad-DOPE (XII).

186 (previously presented).

The method according to claim 168 where the

construct is:

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designated Fucα1-2Galβ1-3(Fucα1-4)GlcNAc-sp-Ad-DOPE (XIII).

187 (previously presented). The method according to claim 168 where the cell or multi-cellular structure is a red blood cell.

188-189 (cancelled).